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What is claimed is:

- 1. A conjugate comprising:
- an Fc portion of an IgA antibody, wherein said Fc portion comprises a CH2 domain and a CH3 domain but does not comprise a hinge region; and

folate or a folate analog.

- 2. The conjugate of claim 1, further comprising a linker.
- 3. The conjugate of claim 2, wherein the linker is a peptide linker.
- **4**. The conjugate of claim **2**, herein the linker is a non-peptide linker.
- 5. The conjugate of claim 1, wherein the Fc portion is a variant Fc.
- **6**. The conjugate of claim **1**, wherein the Fc portion is a variant Fc and the conjugate comprises folate.
- 7. The conjugate of claim 1, wherein the Fc portion is a variant Fc and the conjugate comprises a folate analog.
- **8**. The conjugate of claim **1**, wherein the Fc fragment consists of a CH2/CH3 region from an IgA antibody.
- **9**. The conjugate of claim **1**, formulated for intravenous administration.
- 10. A method of triggering neutrophil (PMN)-mediated apoptosis of folate receptor alpha (FRA)-expressing cancer cells in a subject in need thereof, comprising: administering to the subject a therapeutically effective amount of the conjugate of claim 1, thereby triggering neutrophil (PMN)-mediated apoptosis of folate receptor alpha (FRA)-expressing cancer cells in the subject.
- 11. The method of claim 10, wherein the cancer is breast cancer, ovarian cancer, or lung cancer.
- 12. The method of claim 10, wherein the cancer is triple negative breast cancer.
- 13. The method of claim 10, wherein the cancer includes cancer cells that lack receptors for one or more of estrogen, progesterone and epithelial growth factor.
- 14. The method of claim 10, wherein administration is by intravenous administration.
- 15. A method of recruiting and activating neutrophils to destroy a folate receptor alpha (FRA)-expressing cancer cell in a subject, comprising contacting the cancer cell with a conjugate comprising: